



# Stable carbon isotopes of dissolved inorganic carbon for a zonal transect across the subpolar North Atlantic Ocean in summer 2014

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**Abstract.** The stable carbon isotope composition of dissolved inorganic carbon ( $\delta^{13}\text{C}_{\text{DIC}}$ ) in seawater was measured in samples collected during June–July 2014 in the subpolar North Atlantic. Sample collection was carried out on the RRS *James Clark Ross* cruise JR302, part of the “Radiatively Active Gases from the North Atlantic Region and Climate Change” (RAGNARoCC) research programme. The observed  $\delta^{13}\text{C}_{\text{DIC}}$  values for cruise JR302 fall in a range from  $-0.07$  to  $+1.95\text{‰}$ , relative to the Vienna Pee Dee Belemnite standard. From duplicate samples collected during the cruise, the  $1\sigma$  precision for the 341 results is  $0.08\text{‰}$ , which is similar to our previous work and other studies of this kind. We also performed a cross-over analysis using nearby historical  $\delta^{13}\text{C}_{\text{DIC}}$  data, which indicated that there were no significant systematic offsets between our measurements and previously published results. We also included seawater reference material (RM) produced by A. G. Dickson (Scripps Institution of Oceanography, USA) in every batch of analysis, enabling us to improve upon the calibration and quality-control procedures from a previous study. The  $\delta^{13}\text{C}_{\text{DIC}}$  is consistent within each RM batch, although its value is not certified. We report  $\delta^{13}\text{C}_{\text{DIC}}$  values of  $1.15 \pm 0.03\text{‰}$  and  $1.27 \pm 0.05\text{‰}$  for batches 141 and 144 respectively. Our JR302  $\delta^{13}\text{C}_{\text{DIC}}$  data can be used – along with measurements of other biogeochemical variables – to constrain the processes that control DIC in the interior ocean, in particular the oceanic uptake of anthropogenic carbon dioxide and the biological carbon pump. Our  $\delta^{13}\text{C}_{\text{DIC}}$  results are available from the British Oceanographic Data Centre – doi:10.5285/22235f1a-b7f3-687f-e053-6c86abc0c8a6.

## 1 Introduction

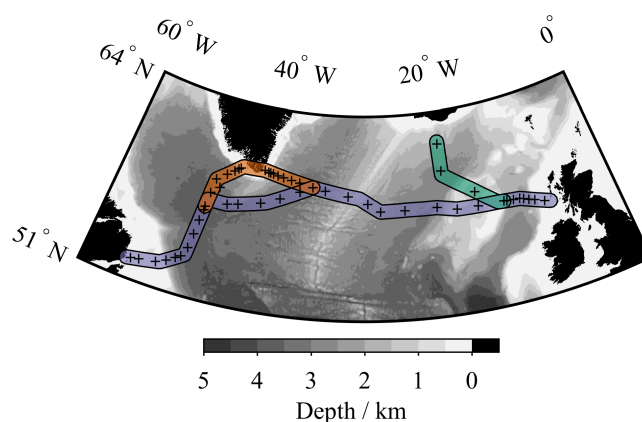
The global ocean has absorbed up to half of the anthropogenic carbon dioxide ( $\text{CO}_2$ ) emitted since the early 1800s (Sabine et al., 2004; Khatiwala et al., 2009, 2013) and it continues to take up about a quarter of annual  $\text{CO}_2$  emissions at the present day (Le Quéré et al., 2009), substantially decreasing  $\text{CO}_2$  accumulation in the atmosphere. The consequences of this uptake include a decline in pH – known as ocean acidification – with lower pH values predicted to persist for cen-

turies longer than the atmospheric  $\text{CO}_2$  anomaly (Caldeira and Wickett, 2003), which will have impacts on marine biogeochemistry and ecology that we are only just beginning to understand (Doney et al., 2009; Achterberg, 2014; Gaylord et al., 2015).

In order to predict the response of the oceanic  $\text{CO}_2$  sink to the continuing rise of the atmospheric partial pressure of  $\text{CO}_2$  ( $p\text{CO}_2$ ), it is useful to first understand the existing spatial distribution of anthropogenic dissolved inorganic carbon ( $\text{DIC}_{\text{anth}}$ ) in the ocean interior. Various methods have

been employed to this end (Sabine and Tanhua, 2010), including back-calculation from DIC, total alkalinity (TA) and dissolved oxygen observations (Brewer, 1978; Chen and Millero, 1979; Gruber et al., 1996); inference from the oceanic distributions of other anthropogenic gases such as chlorofluorocarbons (Hall et al., 2002; Waugh et al., 2006); and multi-linear regressions using measurements from pairs of cruises in the same region, but separated in time (Friis et al., 2005; Tanhua et al., 2007). Oceanic measurements during the past few decades (Quay et al., 2007) and over longer timescales in ice cores (Rubino et al., 2013) show that the rise in  $p\text{CO}_2$  and DIC has been accompanied by a decline in the carbon-13 content of DIC, relative to carbon-12 (reported as  $\delta^{13}\text{C}$ , Eqs. 1 and 2), a phenomenon called the “Suess effect” (Keeling, 1979). This is caused by the lower  $\delta^{13}\text{C}$  of anthropogenic  $\text{CO}_2$  relative to pre-industrial and present-day atmospheric  $\text{CO}_2$ , and it provides another approach to constrain the spatial distribution and inventory of anthropogenic DIC (e.g. Quay et al., 1992, 2003, 2007; Sonnerup et al., 1999, 2007; Körtzinger et al., 2003). The Suess effect has caused significant changes in the present-day distribution of  $\delta^{13}\text{C}_{\text{DIC}}$  in the ocean interior (Olsen and Ninnemann, 2010). Continued observations of oceanic  $\delta^{13}\text{C}_{\text{DIC}}$  are essential for verification of the parameterisations of ocean carbon cycle models (Sonnerup and Quay, 2012).

Here, we present measurements of  $\delta^{13}\text{C}_{\text{DIC}}$  from a zonal transect across the subpolar North Atlantic Ocean in June–July 2014. The cruise, JR302 on the RRS *James Clark Ross*, was carried out as part of the “Radiatively Active Gases from the North Atlantic Region and Climate Change” (RAGNARoCC) research programme. Our observations fill important spatiotemporal gaps in the existing global data set (Schmittner et al., 2013), and will contribute towards the scientific objectives summarised above. Our analysis was carried out following the methodology presented by Humphreys et al. (2015a), but we have been able to make several improvements to the raw data processing and calibration procedures by inclusion of seawater reference material (RM) in every batch of sample analysis. This RM, produced by A. G. Dickson (Scripps Institution of Oceanography, USA), is mainly used for assessing the accuracy of non-isotopic marine carbonate chemistry measurements, and it does not have a certified  $\delta^{13}\text{C}_{\text{DIC}}$  value. Nevertheless, the  $\delta^{13}\text{C}_{\text{DIC}}$  of different RM bottles from the same RM batch should be consistent, allowing us to assess the consistency of our measurements between analysis batches. We determine  $\delta^{13}\text{C}_{\text{DIC}}$  values for the two RM batches that we measured (141 and 144), which could be used to check for systematic offsets between our results and those from other laboratories. We also use the RM results to carry out a statistical analysis of our measurement precision both within and between analysis batches.



**Figure 1.** Bathymetric map of the subpolar North Atlantic Ocean. Black pluses show  $\delta^{13}\text{C}_{\text{DIC}}$  sampling locations during cruise JR302. Coloured sections indicate illustrated transects: blue for Fig. 6, orange for Fig. 7, and green for Fig. 8. Bathymetry data are from the GEBCO\_2014 grid, version 20150318, <http://www.gebco.net>.

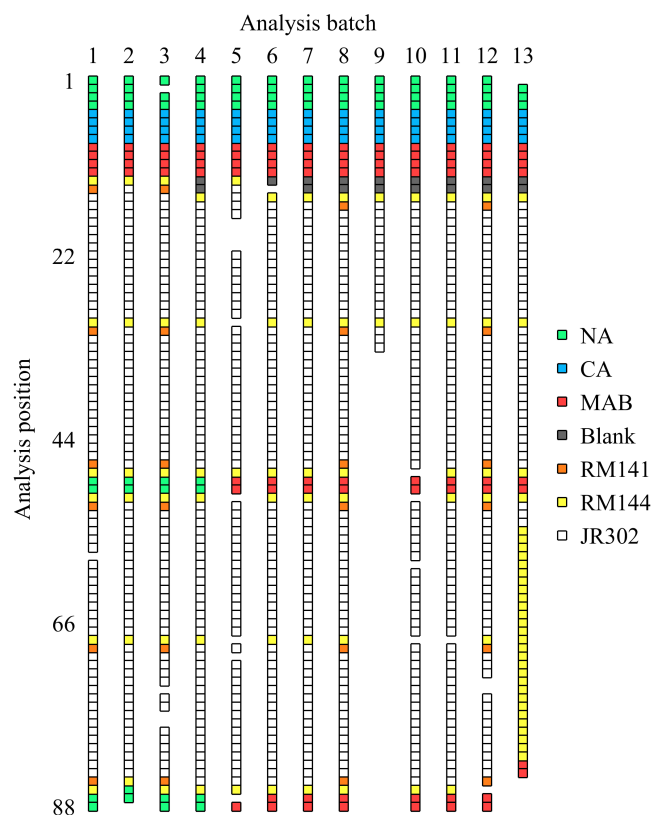
## 2 Sample collection

### 2.1 Cruise details

The  $\delta^{13}\text{C}_{\text{DIC}}$  samples were collected during RRS *James Clark Ross* cruise JR302, which was an approximately zonal transect from St John's, Newfoundland, Canada, to Immingham, UK (Fig. 1), from June to July 2014 (King and Holliday, 2015). During the crossing, several transects were sailed in towards the coast of Greenland, and in the eastern region the ship carried out a short meridional transect north towards Iceland in order to sample the Extended Ellett Line (Holliday and Cunningham, 2013).

### 2.2 Sample collection and storage

Prior to sample collection, the containers were thoroughly rinsed with deionised water (Milli-Q water, Millipore,  $> 18.2 \text{ m}\Omega \text{ cm}^{-1}$ ). Samples were collected from the source (either seawater sampling bottle or underway seawater supply) via silicone tubing, following established best-practice protocols (Dickson et al., 2007; McNichol et al., 2010), as summarised here. The containers were thoroughly rinsed with excess seawater sample immediately before filling until overflowing with seawater, taking care not to generate or trap air bubbles. Two different sample containers were used: (1) 100 mL glass “bottles” with ground glass stoppers, lubricated with Apiezon<sup>®</sup> L grease and held shut with electrical tape, and (2) 50 mL glass “vials”, with plastic screw-cap lids and PTFE/silicone septa. In order to sterilise each sample, 0.02 % of the sample container volume of saturated mercuric chloride solution was added before sealing. A 1 mL air headspace (i.e. 1 % of the sample volume) was also introduced to the bottles, prior to poisoning, by removing this



**Figure 2.** Schematic arrangement of calibration standards (NA, CA and MAB), blanks, RM and samples (JR302) within each analysis batch. Each square represents a separate measurement (i.e. a set of 10 technical replicates). Gaps are left where results have failed quality control.

volume of seawater via pipette. This prevents thermal expansion/contraction of the seawater from breaking the airtight seal. However, the flexible septa on the vials allowed them to be sealed when full of seawater. All samples were stored in the dark until analysis.

### 3 Sample analysis

All of the  $\delta^{13}\text{C}_{\text{DIC}}$  samples were analysed at the Scottish Universities Environmental Research Centre Isotope Community Support Facility (SUERC-ICSF) in East Kilbride (UK), in June–July 2015. We describe the analysis procedure here only in brief, as it was identical to that of Humphreys et al. (2015a).

The samples were analysed in 13 batches. Each analysis batch consisted of up to 88 measurements, 16 of which were of calibration standards, and the remainder were of seawater samples, “blanks” or RM (Fig. 2). Each batch underwent a three-step process of overgassing, equilibration and measurement. For the overgassing step, the air in each of the measurement vials (12 mL Exetainer®) was flushed out and replaced with helium by a PAL system (CTC Analytics). For

equilibration, the standards, samples and RM were reacted with phosphoric acid to convert all DIC to  $\text{CO}_2$ . Finally, the gaseous headspace in each measurement vial was then sampled by the PAL system and transferred to a Thermo Scientific Delta V mass spectrometer via a Thermo Scientific GasBench II, and was measured 10 times (called technical replicates).

For seawater samples and RM, four drops of concentrated phosphoric acid were added to the Exetainer analysis vials prior to overgassing, and 1 mL of liquid sample was then injected into each vial for equilibration. For the standards, only the solid powder standard was added to the analysis vials prior to overgassing, and 1 mL of dilute (10 % by volume) phosphoric acid was added to each vial for equilibration. During batches 4 and 6–13, “blanks” were prepared in the same way as the standards, except that the Exetainer analysis vials were completely empty for overgassing, and had the same dilute acid added as the standards in that batch.

## 4 Measurement processing

We were able to make improvements to the processing of the raw measurements from our previous study (Humphreys et al., 2015a), in part by using the results from the RM that were included in every batch. The processing sequence used in this study is described in full here, including parts that are the same as in Humphreys et al. (2015a); differences between the two approaches are then discussed later. All processing was carried out using MATLAB® (MathWorks, USA).

### 4.1 Definitions

The relative abundance of  $^{13}\text{C}$  to  $^{12}\text{C}$  in a sample  $X$  is given by Eq. (1). The  $R_X$  is then normalised to a reference standard – i.e. Vienna Pee Dee Belemnite (V-PDB) (Coplen, 1995) – using Eq. (2).

$$R_X = \frac{[^{13}\text{C}]_X}{[^{12}\text{C}]_X}, \quad (1)$$

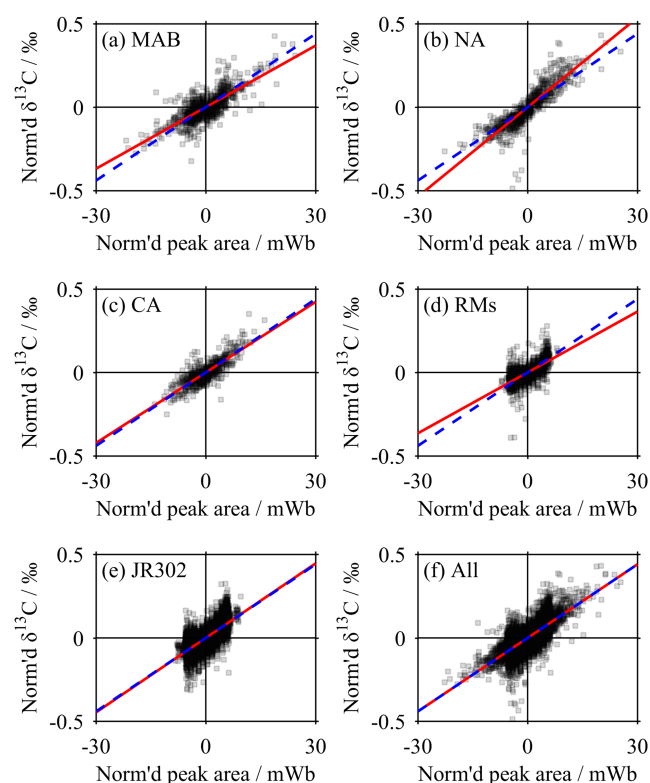
where  $[^{13}\text{C}]_X$  and  $[^{12}\text{C}]_X$  are the concentrations of  $^{13}\text{C}$  and  $^{12}\text{C}$  respectively in  $X$ .

$$\delta^{13}\text{C} = \frac{R_{\text{sample}} - R_{\text{V-PDB}}}{R_{\text{V-PDB}}} \times 1000\text{‰} \quad (2)$$

### 4.2 General procedure

#### 4.2.1 Anomalous measurement removal

Anomalous  $\delta^{13}\text{C}$  measurements were first removed from the sets of technical replicates. These typically occurred when the  $\text{CO}_2$  concentration in a replicate was too low, causing the peak area to fall outside the calibrated range (i.e. the range of peak areas covered by the standards). There were no



**Figure 3.** The peak area correction relationship. All technical replicates are plotted, normalised such that the mean peak area and  $\delta^{13}\text{C}$  for each set of technical replicates are both 0 (‰ or mWb respectively). The red line is the same in each plot, showing the mean relationship used for all of the peak area corrections, while each dashed blue line shows the equivalent relationship determined only from the data scattered in each panel. Individual data points are semi-transparent.

sample measurements with peak areas greater than the calibrated range. Thus all measurements with a peak area less than 5 mWb were judged to be anomalous and discarded, and if this applied to 5 or more of the original 10 technical replicates for a given sample, the entire sample was discarded.

#### 4.2.2 Peak area (linearity) correction

Virtually all samples, standards and RM showed a consistent decline in both peak area and raw  $\delta^{13}\text{C}$  through each set of technical replicates, called “linearity” (Fig. 3). To correct for this, we first “normalised” the peak area and raw  $\delta^{13}\text{C}$  of each set of technical replicates by subtracting the mean peak area and  $\delta^{13}\text{C}$  respectively from each replicate; every set of technical replicates thus had a mean normalised peak area and  $\delta^{13}\text{C}$  of 0 (mWb or ‰ respectively). We then performed an ordinary least-squares linear regression of normalised  $\delta^{13}\text{C}$  against normalised peak area using all technical replicates from all of the samples, standards and RM. The regression was forced through the origin and had a gradient

( $L$ ) of  $0.0147\text{‰ mWb}^{-1}$ . This was used to make a “peak area correction” to all technical replicates:

$$\delta_{\text{lin}} = \delta_{\text{raw}} - L(a - A_{\text{lin}}), \quad (3)$$

where  $\delta_{\text{lin}}$  is the linearity-corrected  $\delta^{13}\text{C}$ ,  $\delta_{\text{raw}}$  is the raw  $\delta^{13}\text{C}$  measurement,  $L$  is the correction gradient (i.e.  $0.0147\text{‰ mWb}^{-1}$ ),  $a$  is the peak area for the technical replicate in mWb, and  $A_{\text{lin}}$  is 20 mWb – the peak area that the correction is made to. The value of  $A_{\text{lin}}$  was chosen because it is the mean peak area for all of the seawater samples, thus minimising the magnitude of this correction.

#### 4.2.3 Averaging

After the peak area correction had been applied, the mean  $\delta^{13}\text{C}$  of each set of technical replicates was calculated. These mean values were then used for the remainder of the data processing.

#### 4.2.4 Blank correction

A “blank correction” was then applied to the standards only (MAB, NA and CA). This was necessary because phosphoric was added to these after the overgassing step, so any  $\text{CO}_2$  dissolved in the acid would be included in the measurement. It was not necessary for the seawater samples and RM, because here the acid was added prior to overgassing. The different procedures were necessary because of the different states of the standards and samples (solid and liquid respectively).

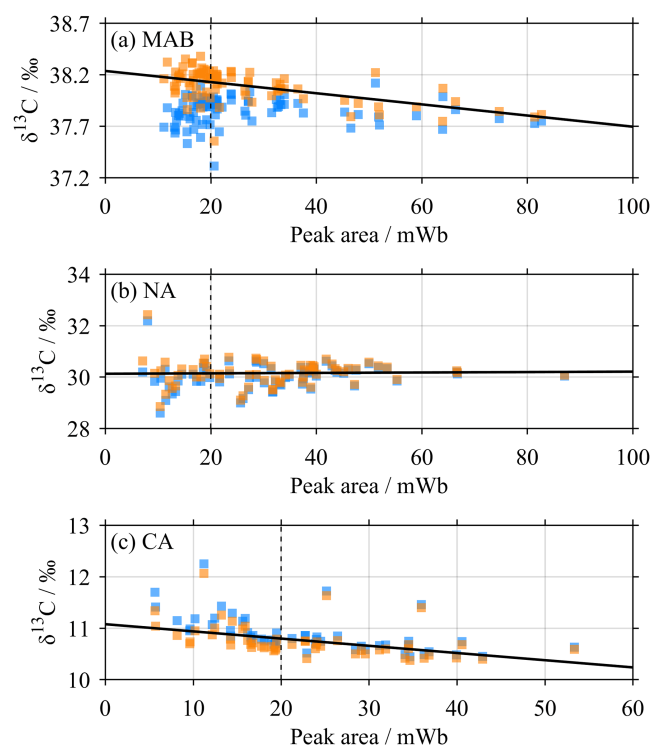
A pair of “blank” measurements were included during analysis batches 4 and 6–13. These were clean, empty Exetainer analysis vials that were otherwise treated in the same way as the standards: acid had been added after overgassing. The mean  $\pm$  standard deviation (SD) peak area and linearity-corrected  $\delta^{13}\text{C}$  of all of these blanks were  $0.277 \pm 0.024$  mWb and  $19.42 \pm 2.47\text{‰}$  respectively. These mean values were then used to make the blank correction to all standards:

$$\delta_{\text{blank}} = \frac{a\delta_{\text{lin}} - A_{\text{blank}}D_{\text{blank}}}{a - A_{\text{blank}}}, \quad (4)$$

where  $\delta_{\text{blank}}$  is the blank-corrected  $\delta^{13}\text{C}$ , and  $A_{\text{blank}}$  and  $D_{\text{blank}}$  are the mean blank peak area and  $\delta^{13}\text{C}$  (i.e.  $0.277 \pm 0.024$  mWb and  $19.42 \pm 2.47\text{‰}$  respectively).

Even after this blank correction, there remained unexplained relationships between peak area and  $\delta^{13}\text{C}$  for the standards (Fig. 4). We therefore performed an ordinary least-squares regression between peak area and blank-corrected  $\delta^{13}\text{C}$  for all measurements in all analysis batches of each standard, and took the value of the regression line at a peak area of 20 mWb as the  $\delta^{13}\text{C}$  value for that standard in order to generate the V-PDB calibration curve (Table 1).





**Figure 4.** The blank correction for the standards. Each panel shows relationships between peak area and  $\delta^{13}\text{C}$  for (a) MAB, (b) NA and (c) CA before (blue) and after (orange) blank correction. The solid black lines are linear least-squares regressions for the data after blank correction. Their intercepts with the dashed black lines at 20 mWb were used as the mean values for each standard to generate the V-PDB calibration curve (Fig. 5). Note that the panels have different axes resolutions.

#### 4.2.5 Calibration to V-PDB

A second-order polynomial fit was generated to determine the certified  $\delta^{13}\text{C}$  values for the standards (Table 1) from their blank-corrected mean values (Fig. 5):

$$\delta_{\text{cert}} = c\delta_{\text{blank}}^2 + s\delta_{\text{blank}} + f, \quad (5)$$

where  $\delta_{\text{cert}}$  is the certified  $\delta^{13}\text{C}$  value (Table 1) and  $c$ ,  $s$  and  $f$  are coefficients describing the curvature, stretch and offset respectively of the calibration fit, taking the following values:  $c = -4.321 \times 10^{-3} \text{‰}^{-1}$ ,  $s = 1.189$ , and  $f = -36.563 \text{‰}$ . Equation (5) was then used to calibrate all of the sample and RM measurements, by inputting the linearity-corrected  $\delta^{13}\text{C}$  values as  $\delta_{\text{blank}}$ ; the output ( $\delta_{\text{cert}}$ ) gives the final, calibrated  $\delta^{13}\text{C}_{\text{DIC}}$ , relative to the V-PDB international standard (Coplen, 1995).

### 4.3 Quality control

#### 4.3.1 Practical problems

We have excluded  $\delta^{13}\text{C}_{\text{DIC}}$  results from our data set where practical problems were encountered and noted during sam-

**Table 1.** The SUERC-ICSF calibration standards.

| Name | Chemical composition | Certified $\delta^{13}\text{C}$ (V-PDB)/‰ | Blank-corrected $\delta^{13}\text{C}$ at 20 mWb/‰ |
|------|----------------------|---|---|
| MAB  | $\text{CaCO}_3$      | +2.48                                     | +38.13  |
| NA   | $\text{NaHCO}_3$     | −4.67                                     | +30.13  |
| CA   | $\text{CaCO}_3$      | −24.23                                    | +10.80  |

ple analysis that discredited specific measurements. For example, during analysis batch 9 the automated needle became detached part way through the overgassing step, resulting in the loss of measurements after this point. The measurements that have been excluded in this way can be identified as gaps in Fig. 2.

#### 4.4 Cross-over analysis

A cross-over analysis was performed using XOVER v1.0.0.1 (Humphreys, 2015) in order to evaluate the consistency of this study's results with “historical” measurements from the Schmittner et al. (2013)  $\delta^{13}\text{C}_{\text{DIC}}$  compilation and our previous study (Humphreys et al., 2014a, b, 2015a). This compilation is probably the best available source for oceanic  $\delta^{13}\text{C}_{\text{DIC}}$  measurements at present, although it may soon be superseded by ongoing efforts to merge and quality-control marine  $\delta^{13}\text{C}_{\text{DIC}}$  data sets (e.g. Becker et al., 2016). The XOVER program follows a similar procedure to the secondary quality control toolbox of Lauvset and Tanhua (2015). Firstly, all historical sampling stations within 150 km of a JR302 CTD station were selected. The 150 km distance is the best compromise for minimising the spatial offset between the JR302 and historical observations while still capturing enough historical data to perform an effective cross-over analysis. At each of these historical stations, a piecewise cubic Hermite interpolating polynomial (PCHIP) fit was generated to predict  $\delta^{13}\text{C}_{\text{DIC}}$  from depth. Values of  $\delta^{13}\text{C}_{\text{DIC}}$  at depths which were equivalent to the JR302 observations were then interpolated using these PCHIP fits. Only JR302 data from deeper than 200 m were used, in order to limit the effect of seasonal variability in  $\delta^{13}\text{C}_{\text{DIC}}$ , which is relatively high near the ocean surface due to biological processes. The differences between the JR302 and historical  $\delta^{13}\text{C}_{\text{DIC}}$  values were calculated and combined into a mean  $\pm$  SD value for each cruise in the historical data sets.

#### 4.5 Precision from duplicates

The SD obtained if one sample was measured many times (i.e.  $1\sigma$  precision, 68.3 % confidence interval) can also be estimated from many duplicate measurements of different samples: it is equal to the mean of the absolute differences between the duplicate pairs divided by  $2/\sqrt{\pi}$  (Thompson and Howarth, 1973; Humphreys et al., 2015a), as follows. For this purpose, each pair of duplicate measurements of a sam-

ple  $i$  is considered to be two values ( $d_{i,1}$  and  $d_{i,2}$ ) that have been randomly selected from a normal distribution with a SD equal to the  $1\sigma$  measurement precision ( $P$ ) and a mean equal to the “true”  $\delta^{13}\text{C}_{\text{DIC}}$  value for that sample. The “duplicate pair difference” ( $D$ ) for the sample  $i$  is then calculated by subtracting the  $\delta^{13}\text{C}_{\text{DIC}}$  of the first duplicate from that of the second (i.e.  $D_i = d_{i,2} - d_{i,1}$ ). As  $P$  is the same for every sample,  $D$  is normally distributed with a mean of 0 and a SD of  $P\sqrt{2}$ . The  $1\sigma$  precision  $P$  can thus be estimated from the SD of all  $D$  by dividing it by  $\sqrt{2}$ . Alternatively, the absolute values of  $D$  follow a half-normal distribution, which has a mean value  $2P/\sqrt{\pi}$ ; thus,  $P$  can also be estimated from the mean of the duplicate pair absolute differences (i.e. the mean of all  $|D|$ ) by dividing the latter by  $2/\sqrt{\pi}$ . This last calculation (Eq. 6) was carried out for all of the analytical duplicate pairs and separately for the sampling duplicates to determine their respective  $1\sigma$  confidence intervals:

$$P = \frac{\sqrt{\pi}}{2N} \sum_{i=1}^N |D_i|, \quad (6)$$

where  $N$  is the total number of duplicate pairs.

## 5 Results and discussion

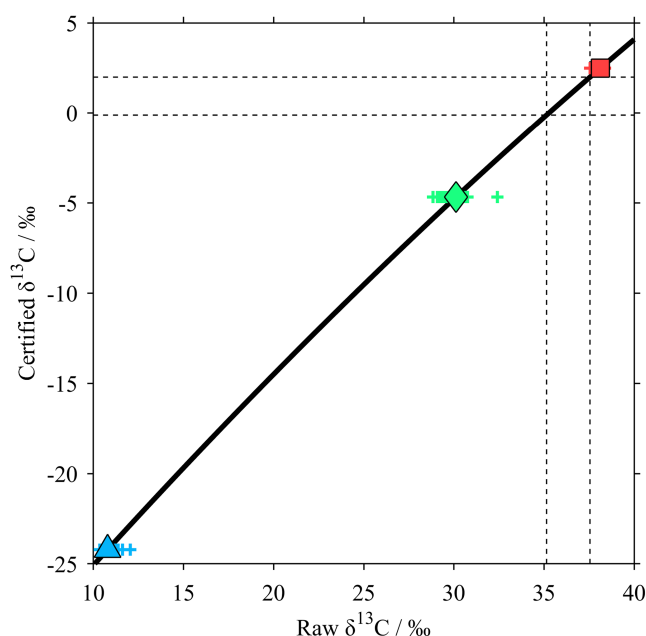
### 5.1 Results in context

#### 5.1.1 Interior $\delta^{13}\text{C}_{\text{DIC}}$ distribution

Our final  $\delta^{13}\text{C}_{\text{DIC}}$  results are presented in Figs. 6–8. To first order,  $\delta^{13}\text{C}_{\text{DIC}}$  is highest in surface waters (shallower than ca. 40 m) taking values up to 2‰. Then,  $\delta^{13}\text{C}_{\text{DIC}}$  decreases with depth to minima just below 0‰ at about 500 m, before increasing again to intermediate values of around 1‰ in deeper waters. This pattern is in general agreement with previous studies (Schmittner et al., 2013).

#### 5.1.2 Cross-over analysis

The cross-over analysis compared the results from this study with three nearby historical cruises: OACES93, 58GS20030922 and D379 (Table 2, Fig. 9). The mean  $\delta^{13}\text{C}_{\text{DIC}}$  residual was significantly different from 0 for all of these cruises ( $p < 0.01$ ). Despite this, the mean ( $\pm$ SD)  $\delta^{13}\text{C}_{\text{DIC}}$  residuals for OACES93 and D379 ( $-0.08 \pm 0.14$ ‰ and  $+0.08 \pm 0.16$ ‰ respectively) were no larger than our reported measurement precision of 0.08‰. Although the mean ( $\pm$ SD) residual for 58GS20030922 was greater ( $-0.19 \pm 0.16$ ‰), it must firstly be considered that this depends on only three matching  $\delta^{13}\text{C}_{\text{DIC}}$  measurements, and secondly that it is still within the range of the accuracy of 0.1–0.2‰ reported for its parent data set (Schmittner et al., 2013). These cruises and JR302 span a time interval of just over 20 years, so invasion of anthropogenic  $\text{CO}_2$  could have modified the  $\delta^{13}\text{C}_{\text{DIC}}$  through the Suess effect (Keeling, 1979), potentially inhibiting the use of cross-over anal-



**Figure 5.** The V-PDB calibration. Mean values for each standard: red square for MAB, green diamond for NA, and blue triangle for CA; results of individual measurements of standards shown as pluses in the same colours as the means. Thick black line shows the calibration curve (Eq. 5); dashed black lines enclose the range of  $\delta^{13}\text{C}$  values for the seawater samples and RM.

ysis. However, in the deeper part of the water column in this region,  $\delta^{13}\text{C}_{\text{DIC}}$  has been observed to change at a rate of less than  $-0.01$ ‰  $\text{yr}^{-1}$  in recent decades (Humphreys et al., 2016). Thus the total change in  $\delta^{13}\text{C}_{\text{DIC}}$  over the period between OACES93 (the earliest historical cruise) and JR302 should be no greater than the stated accuracy of the historical data set, and therefore our cross-over is valid. For future studies this will need to be reconsidered, as it might not hold true over longer timescales. We therefore conclude that any systematic bias between the results of this study and existing  $\delta^{13}\text{C}_{\text{DIC}}$  data sets is negligible relative to the uncertainties of the measurements themselves.

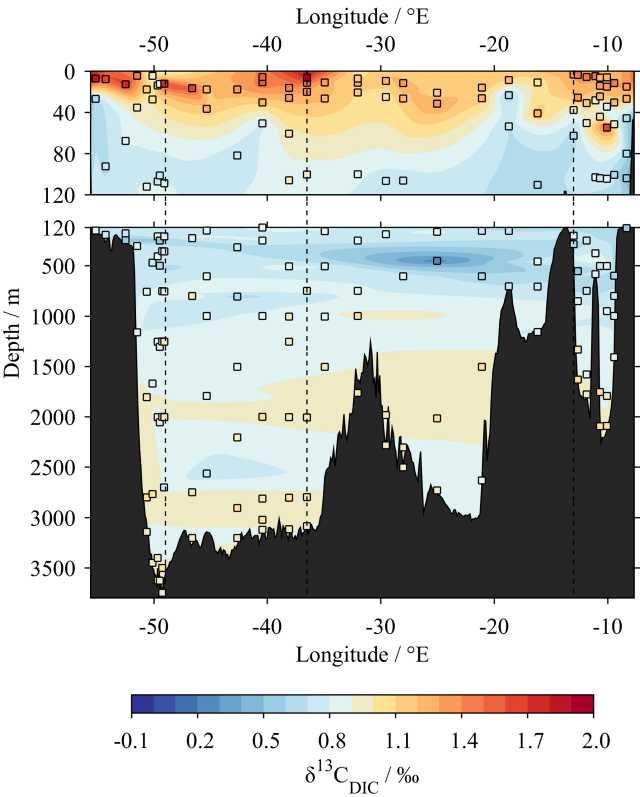
### 5.2 Measurement uncertainty

#### 5.2.1 Sample container types

In tests of duplicate samples collected in these two different container types, Humphreys et al. (2015a) were unable to find evidence of any systematic offset between  $\delta^{13}\text{C}_{\text{DIC}}$  measurements from the same two sample container types that were used in this study. Here, five pairs of sampling duplicates were collected with one sample in each container type. The mean  $\pm$ SD difference in  $\delta^{13}\text{C}_{\text{DIC}}$  for these duplicate pairs was  $-0.01 \pm 0.04$ ‰, with the difference always calculated as the  $\delta^{13}\text{C}_{\text{DIC}}$  value measured in the sample collected in a 50 mL vial subtracted from that collected in a

**Table 2.** Results of the cross-over analysis. Sources: OACES93 on R/V *Malcolm Baldrige*, carbon PIs F. Millero, R. Feely and P. Quay, data from Schmittner et al. (2013); 58GS20030922 on G. O. *Sars*, carbon PIs A. Olsen and T. Johannessen, data from Schmittner et al. (2013); D379 on RRS *Discovery*, carbon PI A. M. Griffiths, data from Humphreys et al. (2015a).

| Cross-over<br>cruise | Sampling<br>date | Mean of $\delta^{13}\text{C}_{\text{DIC}}$<br>residuals/‰ | SD of $\delta^{13}\text{C}_{\text{DIC}}$<br>residuals/‰ | Number of<br>residuals |
|----------------------|------------------|---|---|------------------------|
| OACES93              | Aug 1993         | −0.08   | 0.14  | 19                     |
| 58GS20030922         | Oct 2003         | −0.19   | 0.16  | 3                      |
| D379                 | Aug 2012         | 0.08  | 0.16  | 253                    |

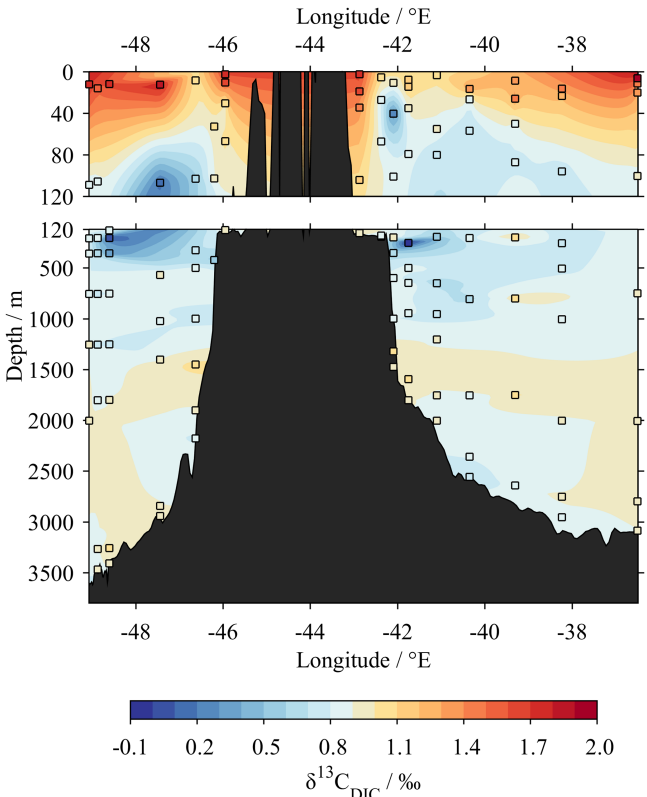


**Figure 6.** Zonal section of  $\delta^{13}\text{C}_{\text{DIC}}$  results from this study, from west to east across the subpolar North Atlantic Ocean (blue in Fig. 1). Coloured squares show actual sample locations and  $\delta^{13}\text{C}_{\text{DIC}}$  values. Vertical dashed lines indicate locations of joints with edges of Figs. 7 and 8. Bathymetry data are from the GEBCO\_2014 grid, version 20150318, <http://www.gebco.net>.

250 mL bottle. A one-sample  $t$  test could not reject the null hypothesis that this mean difference in  $\delta^{13}\text{C}_{\text{DIC}}$  was equal to 0 ( $p = 0.63$ ). We therefore conclude that the container type does not cause a systematic offset in the  $\delta^{13}\text{C}_{\text{DIC}}$  measurement, in agreement with Humphreys et al. (2015a).

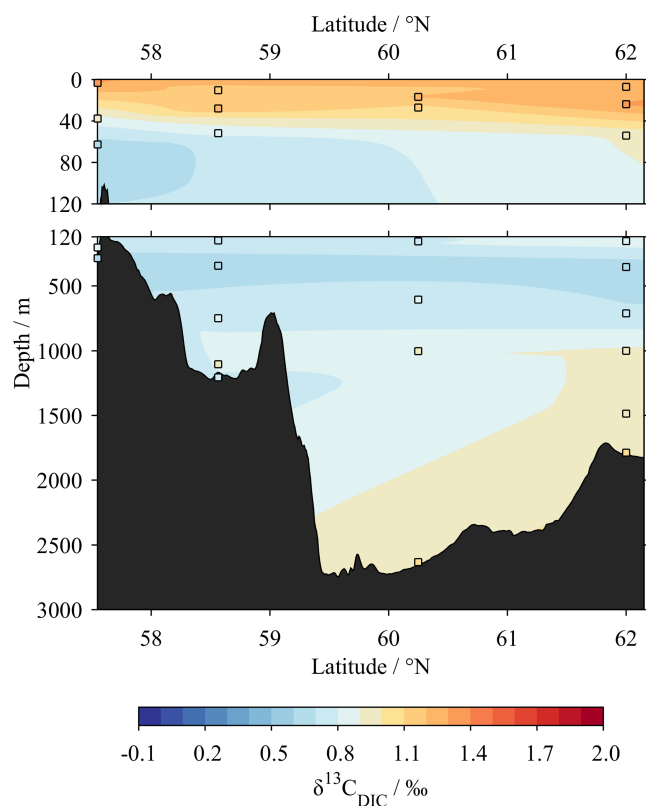
5.2.2 Seawater samples

The typical precision for seawater  $\delta^{13}\text{C}_{\text{DIC}}$  measurements is in the range from about 0.03 to 0.23 ‰ (Olsen et al.,



**Figure 7.** Zonal section of  $\delta^{13}\text{C}_{\text{DIC}}$  results from this study, from west to east near southern Greenland (orange in Fig. 1). Coloured squares show actual sample locations and  $\delta^{13}\text{C}_{\text{DIC}}$  values. Bathymetry data are from the GEBCO\_2014 grid, version 20150318, <http://www.gebco.net>.

2006; Quay et al., 2007; McNichol et al., 2010; Griffith et al., 2012), and we previously reported a value of 0.10 ‰ based on sampling duplicates (Humphreys et al., 2015a). In this study, we again determined the precision of the seawater sample measurements from both analytical and sampling duplicates. There were 341 analytical duplicate pairs, which had a mean absolute difference of 0.075 ‰ and therefore a  $1\sigma$  precision of 0.067 ‰, and 36 sampling duplicate pairs, with a mean absolute difference of 0.090 ‰ and therefore a  $1\sigma$  precision of 0.080 ‰. Although the latter is slightly greater, indicating that the sample collection and storage pro-

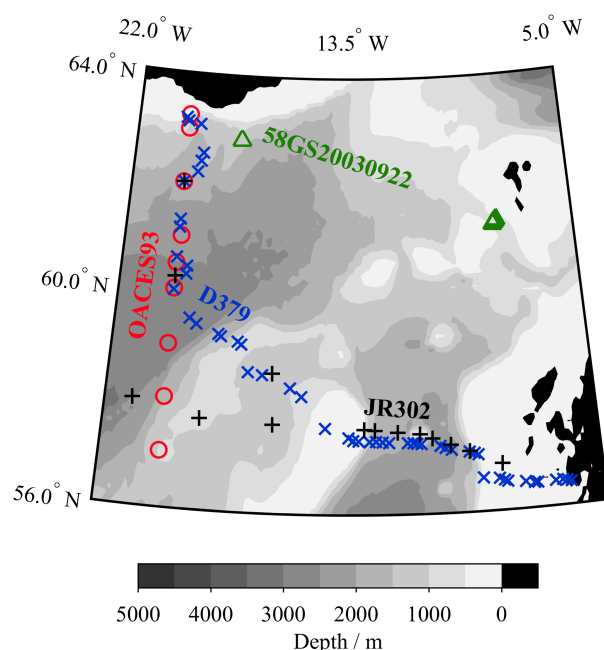


**Figure 8.** Meridional section of  $\delta^{13}\text{C}_{\text{DIC}}$  results from this study, from south to north in the eastern subpolar North Atlantic Ocean (green in Fig. 1). Coloured squares show actual sample locations and  $\delta^{13}\text{C}_{\text{DIC}}$  values. Bathymetry data are from the GEBCO\_2014 grid, version 20150318, <http://www.gebco.net>.

cedures might have adversely affected the measurement precision, Levene's test (Levene, 1960) carried out on the (non-absolute) duplicate differences could not reject the null hypothesis that the analytical and sampling precisions are in fact the same ( $p = 0.33$ ). We therefore report the higher value of  $0.08\text{‰}$  as the  $1\sigma$  precision for this data set; it falls within the range of other studies of this kind. It is important to note that this value is based on consecutively analysed samples, and so might not reflect additional uncertainty engendered by samples being measured non-consecutively or in different analysis batches. However, it is shown in the following section that any such additional uncertainty was negligible.

### 5.2.3 Seawater reference material

Two “batches” of RM were measured in this study: 141 and 144 ([http://cdiac.ornl.gov/oceans/Dickson\\_CRM/batches.html](http://cdiac.ornl.gov/oceans/Dickson_CRM/batches.html)). Each batch consists of multiple bottles of virtually identical seawater. Bottles from within each batch will henceforth be referred to as RM141 and RM144 respectively. The RM are primarily intended for assessment of the accu-



**Figure 9.** Map of JR302 and historical sampling stations used in the cross-over analysis (Table 2). There were no stations suitable for cross-over analysis outside of the area shown on this map. Black pluses show JR302, blue crosses are D379, green triangles are 58GS20030922, and red circles are OACES93. Only historical stations within 150 km of a JR302 station were used, as described in Sect. 4.3.2.

racy of marine carbonate chemistry measurements, in particular DIC and TA (Dickson et al., 2003); they are sterilised and sealed in airtight bottles such that the DIC and TA are consistent in all RM bottles within each RM batch, and so these variables are stable and do not change on timescales of up to a few years. Although the  $\delta^{13}\text{C}_{\text{DIC}}$  value is unknown for these RM, the nature of the preparation and storage process means that we can assume that it is also consistent within each RM batch (A. G. Dickson, personal communication, 18 June 2015), thus allowing us to assess the relative accuracy of our  $\delta^{13}\text{C}_{\text{DIC}}$  measurements between different analysis batches. Unpublished past measurements of  $\delta^{13}\text{C}_{\text{DIC}}$  in similar RM (batches 17, 18 and 19) have supported this assumption, with the  $\delta^{13}\text{C}_{\text{DIC}}$  in multiple (i.e. 3–9) bottles from the same RM batch found to have a SD of about  $0.01\text{‰}$  (A. G. Dickson, personal communication, 18 June 2015).

Typically, we made six measurements of each RM bottle, all within the same analysis batch. These were also spread throughout the analysis batch, and hence not consecutive (Fig. 2). The SD of these results for each bottle therefore represents a longer-term precision estimate than that which we get from the analytical duplicates, which were always analysed one immediately after the other. This approach therefore indicates the reproducibility of measurements carried out anywhere within a single analysis batch (rather than just



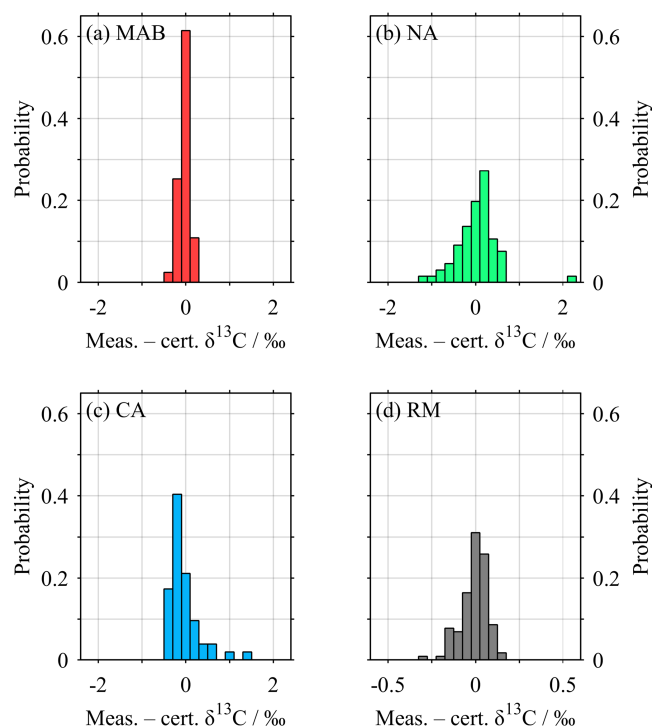
consecutively). The  $1\sigma$  precision from the analytical duplicates was  $0.067\text{‰}$ , while the average SD of the measurements within each RM bottle was slightly smaller ( $0.058\text{‰}$ ). This indicates that the position of samples within each analysis batch did not influence the  $\delta^{13}\text{C}_{\text{DIC}}$  measurement; the relative accuracy of two consecutive measurements is no better than that between measurements from opposite ends of an analysis batch.

The next step is to verify that the difference in  $\delta^{13}\text{C}_{\text{DIC}}$  between different RM bottles of the same batch is negligible. During analysis batch 13, we measured six different RM144 bottles, each up to six times (Table 3). The mean of the six measurements for each RM bottle was taken as its  $\delta^{13}\text{C}_{\text{DIC}}$  value. The SD of these six mean values was  $0.028\text{‰}$ . Next, we compared this to measurements of different RM bottles across different analysis batches. One RM144 bottle was measured during each of analysis batches 1–12. The mean value was calculated for each analysis batch, and the SD of these 12 mean values was  $0.056\text{‰}$ . Although larger than the SD for the six RM bottles within batch 13, this value is still smaller than the overall measurement precision based on samples within the same batch. In addition, we used Levene's test (Levene, 1960) for the null hypothesis that the SD of the 6 RM bottles in analysis batch 13 was the same as the SD of the 12 RM bottles in analysis batches 1–12; the resulting  $p$  value of 0.14 was too great to confidently reject the null hypothesis, so we cannot be certain that there truly is greater variance between analysis batches than within them.

Our final  $\delta^{13}\text{C}_{\text{DIC}}$  values are  $+1.15\text{‰}$  for RM141 and  $+1.27\text{‰}$  for RM144 (Table 3).

#### 5.2.4 Calibration standards

The precision of measurements of the calibration standards was greater (i.e. worse) than that of the samples and RM; all calibrated MAB, NA and CA measurements had SDs of 0.13, 0.46, and  $0.35\text{‰}$  respectively, compared with about  $0.08\text{‰}$  for the RM (Fig. 10). We suggest that this is a result of the necessarily different practical treatment of the liquid seawater samples and RM compared with the powdered solid standards. The former were added to concentrated acid that had been overgassed with helium, while the latter were themselves overgassed prior to addition of dilute acid that may have contained some  $\text{CO}_2$ . The blank correction should have corrected for the influence of this  $\text{CO}_2$ , but there remained an unexplained relationship between peak area and raw  $\delta^{13}\text{C}$  for the standards even after its application (Fig. 4). The poor precision for the standards might be associated with the very small quantities ( $0.1\text{--}1.1\text{ mg}$ ) that were measured out into the Exetainer analysis vials, in contrast to the 1 mL of seawater sample or RM that was used each time – the former would be more susceptible to contamination. This result provides a strong incentive to develop seawater RM with a certified  $\delta^{13}\text{C}_{\text{DIC}}$  value, which can be analysed following the



**Figure 10.** Histograms of offset of all standards and RM from their certified values (Table 1). Mean  $\pm$  SD are (a) MAB,  $-0.03 \pm 0.13\text{‰}$ ; (b) NA,  $+0.02 \pm 0.46\text{‰}$ ; (c) CA,  $-0.04 \pm 0.35\text{‰}$ ; and (d) all RM,  $-0.01 \pm 0.08\text{‰}$ . “Certified” values for RM are our final values,  $1.15\text{‰}$  for RM141 and  $1.27\text{‰}$  for RM144 (Table 3). Note the increased horizontal resolution in (d).

exact same method as the samples during future studies of this kind.

### 5.3 Changes from our previous study

There were three main changes to the data processing from our previous study (Humphreys et al., 2015a): the peak area correction, which previously was carried out using a different relationship for the seawater samples and for each standard; the V-PDB calibration, which was previously carried out separately for each analysis batch; and the drift correction, which is absent in the current study.

#### 5.3.1 Peak area (linearity) correction

The linearity correction in this study was different from that in our previous work (Humphreys et al., 2015a), because we did not find the same relationships between peak area and  $\delta^{13}\text{C}$ . This was due to hardware changes to the mass spectrometer in the intervening time between the studies. We therefore believe that the linearity correction used in each study was appropriate, and would recommend determining the best way to apply this correction on a case-by-case ba-

**Table 3.** Results of the RM measurements. The RM mean and standard deviation (SD) columns contain the mean and SD of the replicate values in each row, except for the rows marked “All”, which contain the mean and SD of all the RM mean values for each RM batch.

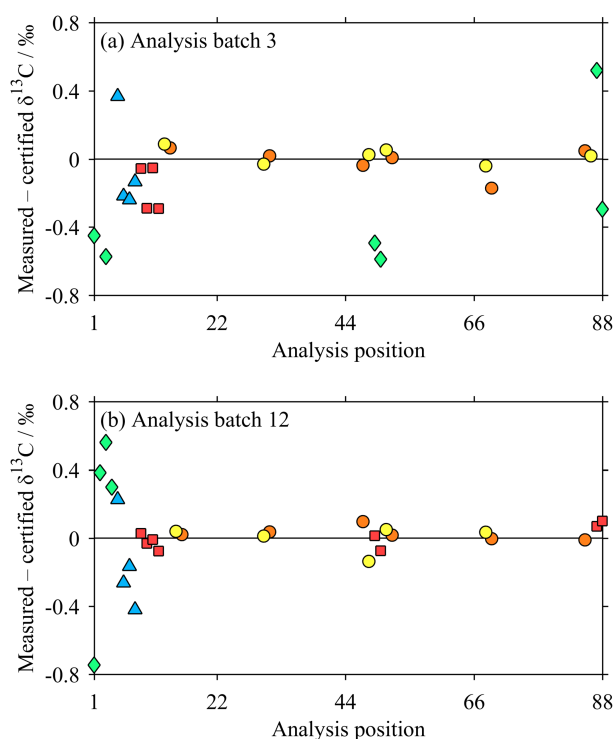
| RM batch | RM bottle | Analysis batch | Replicate RM $\delta^{13}\text{C}_{\text{DIC}}$ measurements/‰ |      |      |      |      |      |  | RM mean/‰ | RM SD/‰ |
|----------|-----------|----------------|--|------|------|------|------|------|--|-----------|---------|
| 141      | 0585      | 1              | 1.03   | 1.10 | 1.09 | 1.10 | 1.21 | 1.13 |  | 1.11      | 0.06    |
| 141      | 0764      | 3              | 1.21   | 1.17 | 1.11 | 1.16 | 0.98 | 1.20 |  | 1.14      | 0.09    |
| 141      | 0455      | 8              | 1.21   | 1.13 | 1.20 | 1.22 | 1.19 | 1.14 |  | 1.18      | 0.04    |
| 141      | 0526      | 12             | 1.17   | 1.19 | 1.25 | 1.17 | 1.15 | 1.14 |  | 1.18      | 0.04    |
| 141      | All       | All            |  |      |      |      |      |      |  | 1.15      | 0.03    |
| 144      | 0030      | 1              | 1.20   | 1.14 | 1.25 | 1.19 | 1.28 | 1.21 |  | 1.21      | 0.05    |
| 144      | 1079      | 2              | 1.20   | 1.22 | 1.32 | 1.28 | 1.27 | 1.12 |  | 1.24      | 0.07    |
| 144      | 1141      | 3              | 1.36   | 1.24 | 1.30 | 1.32 | 1.23 | 1.29 |  | 1.29      | 0.05    |
| 144      | 1024      | 4              | 1.27   | 1.30 | 1.34 | 1.36 | 1.24 | 1.23 |  | 1.29      | 0.05    |
| 144      | 0461      | 5              | 1.29   | 1.24 | 1.18 |      |      |      |  | 1.24      | 0.05    |
| 144      | 0516      | 6              | 1.19   | 1.29 | 1.28 | 1.19 | 1.25 | 1.28 |  | 1.25      | 0.04    |
| 144      | 0399      | 7              | 1.44   | 1.26 | 1.35 | 1.35 | 1.27 | 1.31 |  | 1.33      | 0.06    |
| 144      | 1017      | 8              | 1.12   | 1.16 | 1.10 | 1.07 | 1.18 | 1.10 |  | 1.12      | 0.04    |
| 144      | 0950      | 9              | 1.31   | 1.29 |      |      |      |      |  | 1.30      | 0.01    |
| 144      | 0090      | 10             | 1.27   | 1.33 | 1.36 |      |      |      |  | 1.32      | 0.04    |
| 144      | 0881      | 11             | 1.21   | 1.25 | 1.32 | 1.4  | 1.26 |      |  | 1.29      | 0.07    |
| 144      | 0822      | 12             | 1.31   | 1.28 | 1.13 | 1.32 | 1.30 |      |  | 1.27      | 0.08    |
| 144      | 0151      | 13             | 1.36   | 1.34 | 1.36 | 1.34 | 1.23 |      |  | 1.33      | 0.05    |
| 144      | 0339      | 13             | 1.12   | 1.30 | 1.25 | 1.29 | 1.36 | 1.30 |  | 1.27      | 0.08    |
| 144      | 0575      | 13             | 1.29   | 1.27 | 1.23 | 1.36 | 1.27 |      |  | 1.28      | 0.05    |
| 144      | 0636      | 13             | 1.24   | 1.22 | 1.33 | 1.34 | 1.26 |      |  | 1.28      | 0.05    |
| 144      | 0703      | 13             | 1.17   | 1.29 | 1.25 | 1.33 | 1.30 |      |  | 1.27      | 0.06    |
| 144      | 0745      | 13             | 1.30   | 1.33 | 1.26 | 1.21 | 1.12 |      |  | 1.24      | 0.08    |
| 144      | All       | All            |  |      |      |      |      |      |  | 1.27      | 0.05    |

sis for different data sets. Another result of these hardware changes was a reduction in the mean peak area for the seawater samples from about 35 mWb to about 20 mWb. There is no evidence of any adverse (or particularly beneficial) effects on the quality of the results of either study as a result of these modifications.

### 5.3.2 The V-PDB calibration

The V-PDB calibration in this study was a single equation determined from all of the measurements of all of the calibration standards in every analysis batch, where previously we determined a separate equation for each batch (Humphreys et al., 2015a). This new approach delivered significantly better RM results between batches, as a result of the relatively high uncertainty in the measurements of the calibration standards; the apparent differences in calibration equations between analysis batches were in fact an artefact of these uncertainties. The consequence of this for our previous study is a decrease in precision, but it does not constitute a systematic error. If we apply a different calibration to each batch, as in our previous study, we find mean  $\pm$  SD across all analysis batches of the mean  $\delta^{13}\text{C}_{\text{DIC}}$  of each RM within each batch for RM141 (4 RM bottles across 4 batches) and RM144 (18 RM bottles across 13 batches) of  $1.19 \pm 0.08$  and

$1.30 \pm 0.11$  ‰ respectively; by way of comparison, our new approach in this study yields  $1.15 \pm 0.03$  and  $1.26 \pm 0.05$  ‰ respectively. To determine the significance of these apparent differences, we took the mean RM141 and RM144 results for each analysis batch and used them to test two different null hypotheses, separately for each RM and calibration method. Firstly, we used Welch’s unequal variances *t* test (Welch, 1947) for the null hypothesis that the mean  $\delta^{13}\text{C}_{\text{DIC}}$  across all batches was the same regardless of the V-PDB calibration method. For both RM141 and RM144, the null hypothesis could not be rejected at the 5 % significance level, with *p* values of 0.47 and 0.28 respectively. Secondly, we used Levene’s test (Levene, 1960) for the null hypothesis that the variance of these batch mean results was the same regardless of the V-PDB calibration method. For both RM141 and RM144, the null hypothesis was rejected at the 5 % significance level, with *p* values of 0.01 and 0.03 respectively. Thus we conclude that the change to the V-PDB calibration method – using a single equation across all analysis batches, instead of a separate one for each – results in an improvement in the precision of results from different analysis batches (i.e. reduced SD), and that it does not cause a systematic bias in these results (i.e. no change in the mean).



**Figure 11.** Examples of results of standards and RM within batches (a) 3 and (b) 12. Red squares for MAB, green diamonds for NA, blue triangles for CA, orange circles for RM141, and yellow circles for RM144.

This result cannot necessarily be applied to recalibrate the results of our previous study, as no RM measurements were carried out then. In this study, all measurements were carried out on consecutive days over a 5-week period. However, in the previous study, there were gaps of several weeks between some analysis batches, and there is no way to objectively assess the consistency of the calibration over these breaks retrospectively. Therefore, although the uncertainty estimate for the previous study was probably too generous (i.e. the reported uncertainty was lower than it should have been) and should be approximately doubled, there is no evidence of a systematic bias.

Additionally, the form of the equation used to carry out the V-PDB calibration (Eq. 5) is a second-order polynomial in this study, which differs from the circle used by Humphreys et al. (2015a). This change makes virtually no difference to the final  $\delta^{13}\text{C}_{\text{DIC}}$  results, but it provides a calibration equation that only gives one possible corrected value for each input  $\delta^{13}\text{C}$ . The calibration equation used in this study is also much easier to interpret, with coefficients directly corresponding to the curvature ( $c$ ), stretch ( $s$ ) and translational offset ( $f$ ) of the curve.

### 5.3.3 Drift correction

No drift correction was performed during this study, while Humphreys et al. (2015a) used the measurements of pairs

standards at the middle and end of each analysis batch to correct for instrumental drift. However, the RM measurements spaced throughout each analysis batch in the present study indicated that no drift correction was required, sometimes in disagreement with the calibration standards (Fig. 11). We suggest that this apparent conflict is again a result of the significantly greater uncertainty in individual measurements of the calibration standards relative to those of seawater samples and RM.

As for the V-PDB calibration, this difference does not cause an important systematic offset to the results from our previous study, but rather an increase in the variance. To support this claim, we applied a drift correction following Humphreys et al. (2015a) to our RM144 results in analysis batches 1–8 and 10–12. Analysis batches 9 and 13 were excluded due to the lack of RM data and the different arrangement of RM respectively (Fig. 2). The mean  $\pm$  SD of all individual RM144 measurements (i.e. up to 6 per RM bottle) was  $1.26 \pm 0.08$  ‰ with no drift correction, but  $1.22 \pm 0.17$  ‰ when a drift correction was applied. We used Levene's test (Levene, 1960) to confidently reject the null hypothesis that the SDs were equal with and without the drift correction ( $p = 0.0001$ ), and thus the decline in precision caused by applying the drift correction was significant. We also used Welch's unequal variances  $t$  test (Welch, 1947) for the null hypothesis that the mean value of these RM144 measurements were the same with and without drift correction; although the null hypothesis could be tentatively rejected at the 5 % significance level ( $p = 0.048$ ) the actual magnitude of the difference in the mean values (ca. 0.04 ‰) is smaller than the measurement precision of either study, and can therefore be considered negligible.

However, for the same reasons as given regarding the V-PDB calibration, it would not be appropriate to recommend retrospective changes to the results of our previous study in the absence of any RM measurements therein.

## 6 Conclusions

We successfully measured  $\delta^{13}\text{C}_{\text{DIC}}$  in 341 samples collected from June to July 2014 during RRS *James Clark Ross* cruise JR302 in the subpolar North Atlantic Ocean. The  $\delta^{13}\text{C}_{\text{DIC}}$  values were in the range from  $-0.07$  to  $+1.95$  ‰ relative to V-PDB and had a  $1\sigma$  uncertainty of 0.08 ‰. Our results are internally consistent, with no systematic offsets between or within analysis batches, and a cross-over analysis revealed no systematic bias relative to nearby historical data in deep waters. We have also established  $\delta^{13}\text{C}_{\text{DIC}}$  values for batches 141 ( $+1.15$  ‰) and 144 ( $+1.27$  ‰) of seawater RM obtained from A. G. Dickson (Scripps Institute of Oceanography, USA), and demonstrated that RM bottles within the same batch have consistent  $\delta^{13}\text{C}_{\text{DIC}}$  values. These RMs greatly enhanced our ability to quantitatively assess and improve our data processing approach, and lead us to conclude that the de-

velopment of an internationally available seawater RM with a certified  $\delta^{13}\text{C}_{\text{DIC}}$  value would be a valuable boon to future measurements of this kind.

## 7 Data set availability

The  $\delta^{13}\text{C}_{\text{DIC}}$  measurements described in this study are publicly available, free of charge, from the British Oceanographic Data Centre, with doi:10.5285/22235f1a-b7f3-687f-e053-6c86abc0c8a6 (Humphreys et al., 2015b). The data will also be submitted to the Carbon Dioxide Information Analysis Centre (CDIAC, Oak Ridge National Laboratory, USA) along with other carbonate chemistry and macronutrient metadata from cruise JR302 once those become available.

**Author contributions.** E. Tynan determined the sampling strategy, and E. Tynan, A. M. Griffiths, C. H. Fry and R. Garley collected the samples. F. M. Greatrix, A. McDonald and M. P. Humphreys carried out the measurements and data processing. M. P. Humphreys wrote the manuscript with contributions from all co-authors.

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